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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/891,787	06/26/2001	Carl Nelson Skold		4399

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EXAMINER

DO, PENSEE T

ART UNIT PAPER NUMBER

1641

DATE MAILED: 06/01/2005

Please find below and/or attached an Office communication concerning this application or proceeding.

Office Action Summary

Application No.

09/891,787

Applicant(s)

SKOLD, CARL NELSON

Examiner

Pensee T. Do

Art Unit

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-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☒ Responsive to communication(s) filed on 14 February 2005.
- 2a) ☒ This action is **FINAL**. 2b) ☐ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 54-75 is/are pending in the application.
- 4a) Of the above claim(s) _____ is/are withdrawn from consideration.
- 5) ☐ Claim(s) _____ is/are allowed.
- 6) ☒ Claim(s) 54-72, 74 and 75 is/are rejected.
- 7) ☒ Claim(s) 73 is/are objected to.
- 8) ☐ Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on _____ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some * c) ☐ None of:
- ☐ Certified copies of the priority documents have been received.
 - ☐ Certified copies of the priority documents have been received in Application No. _____.
 - ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- ☒ Notice of References Cited (PTO-892)
- ☐ Notice of Draftsperson's Patent Drawing Review (PTO-948)
- ☐ Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08)
Paper No(s)/Mail Date _____
- ☐ Interview Summary (PTO-413)
Paper No(s)/Mail Date. _____
- ☐ Notice of Informal Patent Application (PTO-152)
- ☐ Other: _____

Conclusion

DETAILED ACTION

Amendment Entry and Claim Status

The amendment filed on February 14, 2005 has been acknowledged and entered.

Claims 54-75 are pending.

Withdrawn Rejection(s)

Rejections under 102 and 103 in the previous office action are withdrawn herein.

NEW GROUNDS OF REJECTION

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

Claims 54-66 are rejected under 35 U.S.C. 103(a) as being unpatentable over Josephson (US 4,672,040) in view of Bradbury et al. (US 5,855,790).

Josephson teaches a method of separating a target material from a liquid mixture, comprising: forming and at least purifying aggregates of two or more crystallites of a magnetizable metal oxide; coating the formed and at least purified aggregates with a polymer such as silage to form coated aggregates and treating the coated aggregates so that the polymer of silane has a binding affinity for the target material to form treated aggregates; combining the treated aggregates with the liquid mixture containing the

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target material for a sufficient time for the target material to bind to the polymer of silane; applying a magnetic field to the combination and separating the treated aggregates, including the target material bound thereto, from the liquid mixture, using the magnetic field. The magnetic particles comprise a magnetic metal oxide core surrounded by an adsorptively or covalently bound silane coat to which a wide variety of bioaffinity adsorbents can be covalently bonded through selected coupling chemistries. The method for preparing the magnetic particles comprises precipitating metal salts in base to form fine magnetic metal oxide crystals. (see col. 9, line 27-col. 13). The target material is an organic material, an organic compound or a biological material such as antibodies, antigens, nucleic acid, etc. The crystallites have a particle size of greater than 500 Angstrom if they are ferromagnetism. The treated aggregates have diameter between 0.1 and 1.5 microns (equivalent to 100 nm – 1500 nm). The magnetizable metal oxide is a magnetizable iron oxide (see col. 12, lines 5-63). The aggregates are formed by a step of treating precipitated magnetite with a base to form a colloidal suspension. The precipitate is washed with sodium chloride solution. (see col. 12, lines 29-33). The treated aggregates are combined with the sample mixture by being dispersed or suspended in a reactor. The aggregates are coated by bonding an organosilane directly to the aggregate of crystallites of the magnetizable metal oxide and bonding the polysaccharide material to the organosilane. (see col. 17, line 50-col. 18, line 25; table III). Bioaffinity adsorbent such as a ligand couples to the organosilane coating of the metal oxide cores by their organofunctionalities. (see col. 7, line 67-col. 8, line 9).

However, Josephson fails to teach the coating is a polysaccharide, coating the aggregates includes steps of bonding the polysaccharide material directly to the aggregate of crystallites of the magnetizable metal oxide;

Bradbury teaches magnetic particles, which comprise a core of a magnetic material, surrounded by a mixture of fibrous material and a solid binding agent. The core consists of particles of iron oxide or other magnetic material. The fibrous material comprises an organic polymer in the form of fibers such as cellulose. Cellulose is a polysaccharide. (see col. 2, lines 43-61; example 2).

It would have been obvious to one of ordinary skills in the art to coat a polysaccharide on the magnetic particles as taught by Bradbury and use in the method of Josephson since both teach the same method of preparing the magnetic particles coated with a polymer. Polymers have the function of protecting the magnetic particles from attack by the aqueous solution. They also contain specific function groups, which are specifically intended to absorb selectively a particular ligand that binds to a substance of interest. With the polymer coating around the magnetic particles, the ligand can be attached to the magnetic particles through the functional groups of the polymer.

Claims 54-66, 68, 69, 74-75 are rejected under 35 U.S.C. 103(a) as being unpatentable over Josephson (US 4,672,040) in view of Kito et al.

Josephson has been discussed above.

However, Josephson fails to teach the coating is a polysaccharide, coating the aggregates includes steps of bonding the polysaccharide material directly to the

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aggregate of crystallines of the magnetizable metal oxide; and a step of aging the crystallites; a step of attaching the polysaccharide material to a pendant functional group on the organosilane; attaching the affinity to the polysaccharide via the functional group; pendant functional group is carboxyl group, a carbonate, an amino group, an aldehyde group, or a sulfonyl group; polysaccharide is a dextran.

Kito teaches a composition containing magnetic metal oxide ultra fine particles which exist in crystal form and comprises an aqueous sol of complex of the magnetic metal oxide ultra fine particles with a polysaccharide, a polysaccharide derivative and/or a protein; and an organic monocarboxylic acid. The magnetic metal oxide ultrafine particles are prepared by: an alkali coprecipitation method, an ion exchange resin method. The alkali coprecipitation method comprises mixing aqueous solution containing divalent metal salt and a trivalent metal salt preferably an iron salt with a base such as NaOH, KOH or NH₄OH; if necessary heating and aging; after separation and water washing of the magnetic metal oxide precipitated, redispersing the magnetic metal oxide in water, and adding a mineral acid such as a hydrochloric acid to obtain a magnetic metal oxide aqueous sol. If necessary, these aqueous sols can be purified and/or concentrated by dialysis, ultrafiltration and centrifugation, etc. The magnetic metal oxide aqueous sol and a polysaccharide and/or protein aqueous solution were mixed to coat the polysaccharide on the magnetic metal oxide (see col. 3, lines 50-col. 4, line 25; col. 8, lines 20-35). The polysaccharide is a dextran. The diameter of the particles is 10 to 500 nm. (see col. 8, lines 57-62). The polysaccharide is a carboxyl polysaccharide, which contains a pendant carboxyl group. (col. 3, line 44). The metal

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salts can be salts with mineral acids such as hydrochloric acid, sulfuric acid, and nitric acid. (see col. 5, lines 23-25).

It would have been obvious to one of ordinary skills in the art to use the idea of coating the polysaccharide as taught by Kito on the magnetic particles such as those in the method of Josephson since both teach coating a polymer on magnetic metal oxide. Coating the polysaccharide on the magnetic particles provides specific functional group that attaches a ligand/binder, which in turn couples to a substance of interest for use in assays. Regarding claim 68, since Josephson teaches that the organosilane couples to the affinity/coupling group via function groups and Kito teaches that polysaccharide has a pendant group, it would have been obvious to one of ordinary skills in the art to couple the affinity/coupling group via the functional group of either the polysaccharide or the organosilane.

Claims 70-72 are rejected under 35 U.S.C. 103(a) as being unpatentable over Josephson and Kito as applied to claims 54-66, 68, 69, 74, 75 above, and further in view of Niswender (US 4, 048,298).

Josephson and Kito have been discussed above.

However, Josephson and Kito fail to teach that the pendant group of the polysaccharide is a carboxyl group attached to the polysaccharide through a linker having at least one heteroatom to every three carbon atoms in the linker; the heteroatom of the linker is oxygen; and the linker is derived from ethylene glycol, an oligoethylene glycol or a polyethylene glycol.

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Niswender teaches a polymeric carrier with a suitable reactive group. The reactive groups are carboxyl, hydroxyl and primary or secondary amine groups. The polymeric material is polysaccharides, dextrin. The reactive group can be crosslinked by inclusion of a substantial amount of a polyethylenically unsaturated monomer, such as ethylene glycol dimethacrylate.(see col. 4, lines 5-45).

It would have been obvious to one of ordinary skills in the art to attach carboxyl group to polysaccharide via an ethylene glycol linker as taught by Niswender to form a polymeric coating on the magnetizable particles of Josephson and Kito since these polymeric coatings are used for attaching ligands/antibody to detect target analyte in assay.

Response to Arguments

Applicant's arguments with respect to claims 54-75 have been considered but are moot in view of the new ground(s) of rejection.

Allowable Subject Matter

Claim 73 is objected to as being dependent upon a rejected base claim, but would be allowable if rewritten in independent form including all of the limitations of the base claim and any intervening claims.

The prior arts fail to teach introducing the pedant functional group to the polysaccharide by reaction with chloroethoxyethoxyacetic acid and base.

Conclusion

Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP

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§ 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

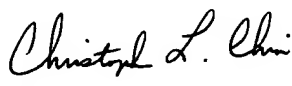
Any inquiry concerning this communication or earlier communications from the examiner should be directed to Pensee T. Do whose telephone number is 571-272-0819. The examiner can normally be reached on Monday-Friday, 7:00-3:00.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Long Le can be reached on 571-272-0823. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Pensee T. Do
Patent Examiner
May 26, 2005


CHRISTOPHER L. CHIN
PRIMARY EXAMINER
GROUP 1800-1641